

Claim 1 line 14 (counting the structural formulae as one line and disregarding the numbering in the left-hand margin), please delete "optionally" and insert --unsubstituted or--.

Claim 8, line 3, please delete "and" (first occurrence) and insert --or a--; line 3 please delete "salts and solvates" and insert --salt or solvate--.

Claim 9, line 16, please delete "and" (first occurrence) and insert --or a--; line 16, please delete "salts and solvates" and insert --salt or solvate--.

Claim 11, line 1, please delete "7" and insert --8--.

Claim 12, line 1, please delete "7" and insert --8--.

Please cancel claims 1.3-21 without prejudice or disclaimer and add the following claims to the application:

(b)

13 --22. A pharmaceutical composition which comprises an effective amount of a compound of formula (I) as defined in claim 1 or a physiologically acceptable salt or solvate thereof together with at least one physiologically acceptable carrier or excipient.--

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--23. A pharmaceutical composition according to claim 22 in a form adapted for oral or parenteral administration.--

--24. A pharmaceutical composition according to claim 22 wherein the active ingredient is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof.--

--25. A pharmaceutical composition according to claim 22 wherein the active ingredient is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one hydrochloride.--

lan H. COATES et al

--26. A method of treating a condition which may be ameliorated by antagonism of 5HT₃ receptors which comprises administering to a patient an effective amount of a compound of formula (I) as defined in claim 1 or a physiologically acceptable salt or solvate thereof to relieve said condition.--

--27. A method according to claim 26 wherein the compound of formula (I) is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof.--

--28. A method according to claim 26 wherein the compound of formula (I) is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one hydrochloride.--

--29. A method according to claim 28 wherein the condition which may be ameliorated by antagonism of 5HT₃ receptors is anxiety.--

--30. A method according to claim 29 wherein the compound of formula (I) is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof.--

--31. A method according to claim 29 wherein the compound of formula (I) is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one hydrochloride.--

-32. A method according to claim 26 wherein the condition which may be ameliorated by antagonism of 5HT₃ receptors is schizophrenia.--

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- --33. A method according to claim 32 wherein the compound of formula (I) is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof.--
- 25 --34. A method according to claim 32 wherein the compound of formula (I) is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1one hydrochloride.--
- --35. A compound according to claim 1 which is 5-ethyl-2,3,4,5-tetrahydro-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof.--
- --36. A method according to claim 26 for the treatment of irritable bowel syndrome.--
- --37. A method according to claim 36 wherein the compound of formula (I) is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof.--
- 27 --38. A method according to claim 36 wherein the compound of formula (I) is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one hydrochloride.--
- --39. A process for the preparation of 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof, which comprises alkylating a compound of formula (II)

